Title 21—Food and Drugs

(This book contains parts 200 to 299)

	Part
CHAPTER I—Food and Drug Administration, Department of Health and Human Services (Continued)	200

CHAPTER I—FOOD AND DRUG ADMINISTRATION, DEPARTMENT OF HEALTH AND HUMAN SERVICES (CONTINUED)

(Parts 200 to 299)

EDITORIAL NOTE: Nomenclature changes to chapter I appear at $59~\mathrm{FR}$ $14366,~\mathrm{Mar}.$ $28,~1994,~\mathrm{and}$ $66~\mathrm{FR}$ $56035,~\mathrm{Nov}.$ 6,~2001.

SUBCHAPTER C—DRUGS: GENERAL

Part	
200	General
201	Labeling
202	Prescription drug advertising
203	Prescription drug marketing
205	Guidelines for State licensing of wholesale pre- scription drug distributors
206	Imprinting of solid oral dosage form drug products for human use
207	Registration of producers of drugs and listing of drugs in commercial distribution
208	Medication Guides for prescription drug products
210	Current good manufacturing practice in manufacturing, processing, packing, or holding of drugs; general
211	Current good manufacturing practice for finished pharmaceuticals
216	Pharmacy compounding
225	Current good manufacturing practice for medicated feeds
226	Current good manufacturing practice for Type A medicated articles
250	Special requirements for specific human drugs
290	Controlled drugs
299	Drugs; official names and established names